

44114–2600, at the Region 5 Office of the Environmental Protection Agency, 77 West Jackson Street, Chicago, Illinois 60604–3590, and at the Consent Decree Library, 1120 G Street, NW., 4th Floor, Washington, DC 20005, (202) 624–0892. A copy of the proposed Consent Decree may be obtained in person or by mail from the Consent Decree Library, 1120 G Street, NW., 4th Floor, Washington, DC 20005. In requesting a copy, please enclose a check in the amount of \$19.75 (25 cents per page reproduction cost) payable to the Consent Decree Library.

**Joel M. Gross,**

*Chief, Environmental Enforcement Section, Environment and Natural Resources Division.*

[FR Doc. 98–18901 Filed 7–15–98; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importation of Controlled Substances; Notice of Application

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on May 19, 1998, Knight Seed Company, Inc., 151 W. 126th Street, Burnsville, Minnesota 55337, made application by renewal to the Drug Enforcement Administration to be registered as an importer of marihuana (7360), a basic class of controlled substance listed in Schedule I.

This application is exclusively for the importation of marihuana seed which will be rendered non-viable and used as bird seed.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance may file written comments on or objections to the application described above, and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing may be addressed,

in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (30 days from publication).

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import a basic class of any controlled substance in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1301.34(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: July 2, 1998.

**John H. King,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

[FR Doc. 98–18894 Filed 7–15–98; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated January 21, 1998, and published in the **Federal Register** on February 12, 1998, (63 FR 7181), Knoll Pharmaceutical Company, 30 North Jefferson Road, Whippany, New Jersey 07981, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Dihydromorphine (9145) .....	I
Hydromorphone (9150) .....	II

The firm plans to produce bulk product and finished dosage units for distribution to its customers.

DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Knoll Pharmaceutical Company to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that

the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: June 30, 1996.

**John H. King,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

[FR Doc. 98–18895 Filed 7–15–98; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Office of Justice Programs

[OJP (OJP)–1185]

RIN 1121–ZB22

#### State and Local Domestic Preparedness Equipment Support Program

**AGENCY:** Office of Justice Programs, Office for State and Local Domestic Preparedness Support (OSLDPS), Justice.

**ACTION:** Request for proposals.

**SUMMARY:** The Office for State and Local Domestic Preparedness Support is soliciting grant applications from Chief Executive Officers (CEO) in targeted jurisdictions; e.g., selected counties and cities, to fund the acquisition of certain types of equipment in the following categories: personal protective equipment, and detection, decontamination, and communications equipment. This equipment will be needed by first responders; i.e., fire services, emergency medical services, hazardous materials response units, and law enforcement agencies, to respond to a terrorist incident and the use of weapons of mass destruction.

**DATES:** Applications for funding must be received by the Office for State and Local Domestic Preparedness Support not later than July 17, 1998.

**ADDRESSES:** Applications must be mailed to: Office for State and Local Domestic Preparedness Support, 810 7th St., NW, Washington, D.C. 20531.

**FOR FURTHER INFORMATION CONTACT:** The National Criminal Justice Reference Service (NCJRS) at 1–800–688–4252 or the U.S. Department of Justice Response Center at 1–800–421–6770.

#### SUPPLEMENTARY INFORMATION:

##### Authority

This action is authorized under Public Law 105–119; the Departments of Commerce, Justice, and State; the